IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of: Robert Hugh BRADBURY et al.)) Confirmation No.: 3653			
Application No.: 10/578,663)	Group Art Unit: Unassigned			
Filed: May 9, 2006)	Examiner: Unassigned			
For: QUINAZOLINE DERIVATIVES)	Date: January 17, 2007			
Commissioner for Patents U.S. Patent and Trademark Office					
Customer Window, Mail Stop Amendment					
Randolph Building					
401 Dulany Street					

Sir:

Alexandria, VA 22314

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

UNDER 37 C.F.R. § 1.97(b)

Citation of Technically Related U.S. Patent Applications

The Examiner's attention is directed to the following technically related U.S. patents or applications of Applicants' assignee:

Inventor	U.S. Serial No.	U.S. Patent No.	PCT Publication No.	
	Filing Date	Issue Date	PCT Publication Date	
Hennequin et al.	11/443,395		WO 03/040109	
	May 31, 2006		May 15, 2003	
Hennequin et al.	11/443,208		WO 03/040108	
	May 31, 2006		May 15, 2003	
Bradbury et al.	10/554,202		WO 2004/093880	
	October 24, 2005		November 4, 2004	
Bradbury et al.	10/571,851		WO 2005/026151	
	March 15, 2006		March 24, 2005	
Bradbury et al.	10/572,262		WO 2005/026152	
	March 16, 2006		March 24, 2005	
Bradbury et al.	10/857,342		WO 2005/012290	
	June 1, 2004		November 4, 2004	
Bradbury et al.	11/628,011		WO 2005/118572	
	November 30, 2006	<u>L</u>	December 15, 2005	

With the exception of the documents listed in the table above in bold (for which a copy was provided with the previously filed Information Disclosure Statement), a copy of the specification and claims for each application, in the form of the published PCT application from which such application was filed is being filed herewith. Consideration of each listed application is earnestly solicited

Pursuant to 37 C.F.R. §§ 1.56 and 1.97(b), Applicants request the Examiner to consider this Supplemental Information Disclosure Statement and the documents listed on the attached Form PTO-1449. To the best of the undersigned's knowledge, this Supplemental Information Disclosure Statement is being filed before the mailing date of a first Office Action on the merits for the above-referenced application. Accordingly, Applicants do not believe a fee is due for filing this Supplemental Information Disclosure Statement.

Copies of the listed documents are attached. Applicants respectfully request that the Examiner initial and return the Form PTO-1449, indicating that the information has been considered and made of record herein.

This submission does not represent that a search has been made or that no better art exists and does not constitute an admission that each or all of the listed documents are material or constitute "prior art." Applicants reserve the right to take appropriate action to establish the patentability of the disclosed invention over the listed documents, should one or more of the documents be applied against the claims of the present application.

Except for issue fees payable under 37 C.F.R. §1.18, the Commissioner is hereby authorized by this paper to charge any additional fees during the entire pendency of this application including fees due under 37 C.F.R. §§1.16 and 1.17 which may be required, including any required extension of time fees, or credit any overpayment to Deposit Account No. 50-0310. This paragraph is intended to be a

CONSTRUCTIVE PETITION FOR EXTENSION OF TIME in accordance with 37 C.F.R.

§1.136(a)(3).

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By:

January 17, 2007 Date:

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INFORMATION DISCLOSURE CITATION		Attorney Docket No. 056291-5283			Application No. 10/578,663					
(Use several sheets if necessary)			Applicants: Robert Hugh BRADBURY et al.							
PTO Form 1449 January 17, 2007			Filing Date: May 9, 2006		006	Group Art Unit: Unassigned				
			U.S. PA	TENT DO	CUMENTS					
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	1	Document No.	POREIGN Date	PATENT	DOCUMENTS Country	Class	Sub-Class	Translation		
	1.	WO 2005/012290	November 4, 200	4	WIPO					
	2.	WO 2005/118572	December 15, 200		WIPO					
	3.	WO 2006/008526	January 26, 2006	5	WIPO					
										
		OTHER DOC	CUMENTS (Includ	ding Auth	or, Title, Date, Pe	rtinent Pa	ges, etc.)			
	4.	Ballard et al. "Inhibitors				nase: optim	isation of po	tency and in vivo		
	_	pharmacokinetics" Bioo	rg Med Chem Lett.	16(18):49	08-4912 (2006)		C41	line anne" Tetrologia		
	5.	Harris et al. "Facile synthesis of 7-amino anilinoquinazolines via direct amination of the quinazoline core" Tetrahedron letters 46(43):7381-7384 (2005)								
	6.			ihvdroxva	uinazoline" Tetrah	edron lette	rs 46(45):77	15-7719 (2005)		
	 6. Harris et al. "Selective alkylation of a 6,7-dihydroxyquinazoline" Tetrahedron letters 46(45):7715-7 7. Hennequin et al. "Novel 4-anilinoquinazolines with C-6 carbon-linked side chains: synthesis and str 									
		relationship of a series of potent, orally active, EGF receptor tyrosine kinase inhibitors" Bioorg Med Chem Lett. 16(10):2672-2676 (2006)								
	8.	Stamos et al. "Structure of the Epidermal Growth Factor Receptor Kinase Domain Alone and in Complex with a 4-Anilinoquinazoline Inhibitor" J. Biol. Chem. 277(48):46265-46272 (2002)								
	9.			inhibitors in cancer treatment" Exp. Opin. Ther. Patents 7(6):571-588 (1997)						
	10.	Traxler et al. "Tyrosine kinase inhibitors in cancer treatment (Part II)" Exp. Opin. Ther. Patents 8(12):1599-1625 (1998)								
	11.	Tsou et al. "6-Substituted-4-(3-bromophenylamino)quinazolines as Putative Irreversible Inhibitors of the Epidermal Growth Factor Receptor (EGFR) and Human Epidermal Growth Factor Receptor (HER-2) Tyrosine Kinases with Enhanced Antitumor Activity" J. Med. Chem. 44:2719-2734 (2001)								
	12.	Vema et al. "Design of EGFR kinase inhibitors: a ligand-based approach and its confirmation with structure-based studies" Bioorg Med Chem. 11(21):4643-4653 (2003)								
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Examiner	<u> </u>	1	Date C	Considered		******				
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